

(19) World Intellectual Property Organization
International Bureau



(43) International Publication Date
17 October 2002 (17.10.2002)

PCT

(10) International Publication Number
WO 02/081366 A1

(51) International Patent Classification⁷: **B82B 3/00**, 1/00, C01B 31/02, C23C 16/46, D01F 9/12, 9/127, 9/133

(21) International Application Number: PCT/AU02/00437

(22) International Filing Date: 4 April 2002 (04.04.2002)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:
PR 4217 4 April 2001 (04.04.2001) AU

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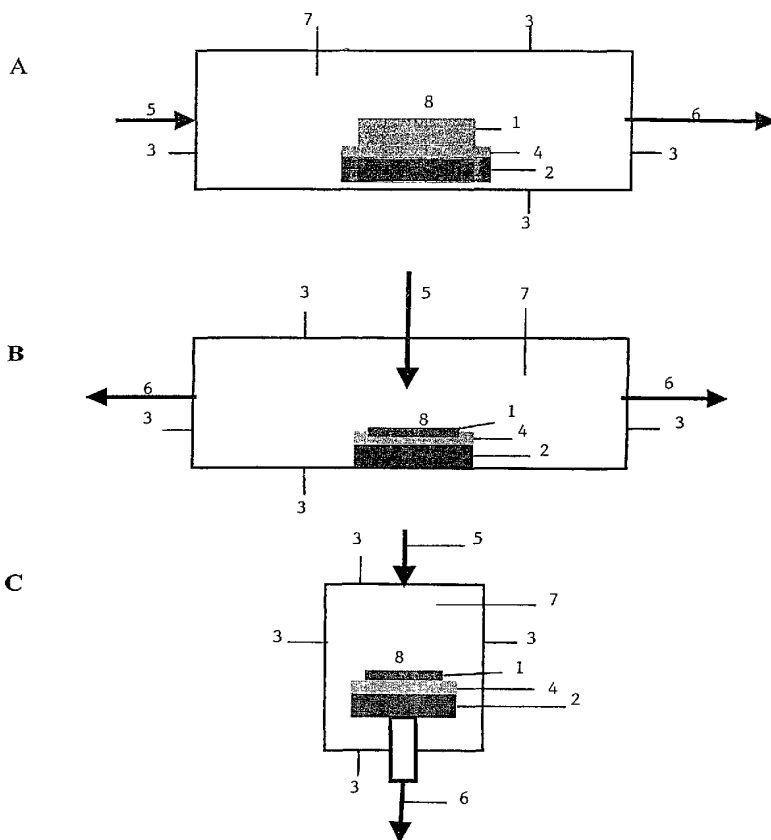
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(81) Designated States (national): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,

[Continued on next page]

(54) Title: PROCESS AND APPARATUS FOR THE PRODUCTION OF CARBON NANOTUBES



(57) Abstract: A process for preparing carbon nanotubes comprising locating a substrate (1) capable of supporting carbon nanotube growth in a localised heating zone within a reaction chamber (7), said localised heating zone being provided by a heating element (2) located within the reaction chamber (7), passing a gaseous carbonaceous material into the reaction chamber such that the gaseous material passes over and contacts the substrate (1) in the localised heating zone, whereby the gaseous material undergoes pyrolysis under the influence of the heat to form carbon nanotubes on the substrate (1). Embodiments of the process prepare multilayer carbon nanotubes and hetero-structured multilayer carbon nanotube films.



WO 02/081366 A1



GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW.

GB, GR, IE, IT, LU, MC, NL, PT, SE, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

Published:

— with international search report

(84) Designated States (regional): ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR,

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

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Process and Apparatus for the Production of Carbon Nanotubes

This invention relates to carbon nanotubes, in particular to a process and apparatus for the preparation of carbon nanotubes.

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Carbon nanotubes usually have a diameter in the order of 0.4 nanometres to 100 nanometres and a length of up to about 1 centimetre. These elongated nanotubes consist of carbon hexagons arranged in a concentric manner with both ends of the tubes normally capped by pentagon-containing fullerene-like structures. Carbon nanotubes may have a single wall or multiwall structure. They can behave as a semiconductor or metal depending on their diameter and helicity of the arrangement of graphitic rings in the walls, and dissimilar carbon nanotubes may be joined together allowing the formation of molecular wires with interesting electrical, magnetic, nonlinear optical, thermal and mechanical properties. These unusual properties have led to diverse potential applications for carbon nanotubes in material science and nanotechnology. Indeed, carbon nanotubes have been proposed as new materials for electron field emitters in panel displays, single-molecular transistors, scanning probe microscope tips, gas and electrochemical energy storages, catalyst and proteins/DNA supports, molecular-filtration membranes, and energy-absorbing materials (see, for example: M. Dresselhaus, *et al.*, *Phys. World*, January, 33, 1998; P.M. Ajayan, and T.W. Ebbesen, *Rep. Prog. Phys.*, 60, 1027, 1997; R. Dagani, *C&E News*, January 11, 31, 1999). The importance of carbon nanotechnology is evidenced by increasing research and development funding (*C & E News*, May 1, 2000, pp.41-47).

For most of the above applications, it is highly desirable that the carbon nanotubes are aligned and/or formed into patterns so that the properties of the individual nanotubes can be easily assessed and they can be incorporated effectively into devices.

Carbon nanotubes have been synthesised using arc discharge (S. Iijima, *Nature*, 354, 56-68, 1991; T. W. Ebbesen and P. M. Aegean, *Nature*, 358, 220-222, 1992) and catalytic pyrolysis (see, for example: M. End. *et Al J. Pays Chum. Solids*, 54, 1841-1848, 1994; V. Ivanov, *et Al Chum. Pays. Let.* 223, 329-335, 1994) and often exist in an randomly

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- entangled state. Patterned and non-patterned carbon nanotube films having the nanotubes aligned perpendicularly with the substrate have been prepared by pyrolysis of iron (II) phthalocyanine in a flow reactor comprising a quartz glass tube heated by a dual furnace (*J. Phys. Chem. B.*, **104**, 2000, 1891). Ren *et al.*, have synthesised large arrays of well-aligned carbon nanotubes by radio-frequency sputter-coating of a thin nickel layer onto a substrate, followed by plasma-enhanced hot filament chemical vapour deposition of acetylene in the presence of ammonia gas at approximately 666°C (*Science*, **282**, 1998, 1105).
- 10 Carbon nanotubes may be prepared at a variety of temperatures, although generally higher temperatures, for example, 600°C to 1100 °C, are required for the preparation of aligned carbon nanotubes. For economic reasons it is preferable to prepare carbon nanotubes at lower temperatures, for example, between 300°C to 800 °C.
- 15 Carbon nanotubes can be prepared in flow reactors comprising a glass tube surrounded by a dual furnace. This technique results in the entire reactor, including the glass tube, being heated and maintained at pyrolysis temperature. Furthermore, carbon is not only deposited on the substrate, but also on the other hot surfaces in the reactor, such as the inside of the glass tube. The carbon deposits on the glass can obscure the view of the substrate, making it difficult to visually monitor the growth of the nanotubes. The positioning of the furnace also generally obscures the view of the substrate and the growth of the nanotubes.
- 20

It is an object of the present invention to overcome or at least alleviate one or more of the disadvantages of the prior art.

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According to a first aspect of the invention there is provided a process for preparing carbon nanotubes comprising:

- 30 locating a substrate capable of supporting carbon nanotube growth in a localised heating zone within a reaction chamber, said localised heating zone being provided by a heating element located within said reaction chamber,

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passing a gaseous carbonaceous material into said reaction chamber such that the gaseous material passes over and contacts said substrate in the localised heating zone, whereby said gaseous material undergoes pyrolysis under the influence of
5 said heat to form carbon nanotubes on said substrate.

According to a second aspect of the invention there is provided a reactor for preparing carbon nanotubes comprising:

10 a reaction chamber,

at least one support means located within said reaction chamber capable of supporting a substrate, said substrate being capable of supporting carbon nanotube growth,

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at least one heating element located within said reaction chamber capable of providing localised heating to said substrate within said reaction chamber,

means for passing a gaseous carbonaceous material into said reaction chamber such
20 that it passes over and contacts said substrate.

According to the present invention the substrate is heated by a heating element in a localised heating zone within a reaction chamber, thereby avoiding the need to heat the entire reaction chamber to pyrolysis temperatures. While the pyrolysis can be achieved at
25 any suitable temperature in the localised heating zone, the process of the invention conveniently allows the preparation of carbon nanotubes at temperatures as low as 300°C. The carbon nanotubes are grown on a substrate that is heated to the required temperature by the heating element. In view of the lower temperatures required and the fact that the heating is localised, the present invention can provide substantial energy and cost savings
30 relative to conventional methods. Also, since the heating is localised to the heating zone, the growth of carbon nanotubes at sites within the reaction chamber other than on the

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substrate and the production of amorphous carbon byproducts inside the reaction chamber are minimised. This also leads to a cleaner reaction chamber and purer carbon nanotube films being formed. If amorphous carbon is deposited on other hot surfaces, for example, exposed areas of the heating element, they are readily removed by heating the heating
5 element in air, causing the amorphous carbon to be oxidised to CO₂. The reaction chamber therefore may be easily cleaned.

The reaction chamber may be defined by one or more walls, and may be of any size or shape suitable for accommodating one or more heating elements. The wall(s) may be
10 formed from any suitable material, including metals, such as steel, aluminium, copper, silver, platinum or alloys, glass, such as quartz glass, normal glass or the like, plastic, polymethylmethacrylate (PMMA), Mylar, polypropylene (PP), polyethylene (PE) or their composites, or a combination thereof. The localised heating zone in the vicinity of the heating element ensures that the temperature of the wall(s) of the chamber remain lower
15 than the temperature in the localised heating zone where pyrolysis occurs. If the chamber is large enough, walls of the chamber remote from the heating zone will remain at ambient temperature. Preferably the reaction chamber is formed from glass or at least includes one or more glass panels. Preferably, the chamber is tubular and its walls are transparent or partially transparent. Advantageously, transparency allows the visual observation of
20 carbon nanotube growth and therefore undesirable growth can be terminated at any stage. Visual observation also allows easier control of the length of the carbon nanotubes by stopping the growth process at a desired time.

The support means may be any support means capable of supporting a substrate within the
25 reaction chamber and capable of withstanding the pyrolysis temperatures used. For example, the support means may be in the form of a solid block, plate, grate, bracket, cradle, stretcher, scaffold or the like and may be made from any suitable material, for example, metal or ceramic materials. The support means may be any size or shape suitable to support the substrate.

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The heating element may be any suitable heating means capable of heating a substrate and

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providing a localised heating zone. For example, suitable heating means may include resistant wires, induction field, microwave radiation or infrared radiation. The localised heating zone can also be heated from a remote point by, for example, a focussed infrared beam or laser beam. In a preferred embodiment, the heating element also acts as the support means for the substrate. In this embodiment, the heating element preferably forms a flat surface upon which the substrate is supported. An example of a suitable heating element which also acts as a substrate support is a ceramic plate into which resistant wires have been inserted. The heating element/substrate support may be formed in any shape or size appropriate to support and heat the substrate and heating zone. Preferably, the heating element allows the substrate to be heated homogenously, i.e., the temperature distribution of the heated substrate is homogenous. One means of achieving homogenous temperature distribution is to place a conducting material, for example, a copper sheet, between the heating plate and the substrate allowing even temperature distribution.

The reactor of the invention also includes a means for passing gaseous carbonaceous material into the reaction chamber such that it passes over and contacts the substrate. This means may be provided by at least one gas conduit. The at least one gas conduit is positioned to allow the flow of gaseous carbonaceous material into the localised heating zone. In a preferred embodiment, the inlet for the carbonaceous material is positioned directly above the substrate so that the gaseous carbonaceous material is supplied directly to the localised heating zone. Alternatively, the gaseous carbonaceous material may be supplied through an inlet at one end of the chamber and allowed to flow across the substrate in the localised heating zone. Multiple gas conduits may be used to supply gaseous carbonaceous material to a large localised heating zone or multiple localised heating zones located within the reaction chamber.

The at least one gas conduit may also be used as a gas inlet for supplying other gases to the reaction chamber and as a gas outlet to allow the exit of gases from the chamber. One gas conduit may be used as both gas inlet and gas outlet. Alternatively, multiple gas conduits may be used, each functioning as a gas inlet or gas outlet.

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If a single gas conduit is used, it may be attached to all gas sources to be supplied to the chamber and a vacuum so the chamber may be evacuated. However, the vacuum may not be applied to the chamber at the same time as gas is supplied. The vacuum is not necessary if inert gases (e.g. Ar) are used to flush the reaction chamber.

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A gas inlet may be used to supply reducing or inert atmospheres, for example, H₂ and/or nitrogen or argon, to the chamber before pyrolysis and to supply the gaseous carbonaceous material to be pyrolysed. These gases may be supplied through a single inlet or through separate inlets.

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A gas outlet may be used to allow the exit of the unused gases and byproducts of the pyrolysis reaction. A gas outlet may be attached to a vacuum pump to allow evacuation of the reactor before the introduction of a reducing and/or inert atmosphere. The gas outlet may also be attached to a device, such as a bubbler, to allow a slight positive pressure of gas to be maintained in the chamber during the deposition of carbon nanotubes.

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In the process of the invention, the substrate may be any substrate capable of withstanding the pyrolysis conditions employed and capable of supporting carbon nanotube growth. Examples of suitable substrates include quartz glass, mesoporous silica, nanoporous alumina, ceramic plates, glass, graphite and mica. Preferably the substrate is ordinary glass. Preferably the surface of the substrate upon which the carbon nanotubes are grown is smooth.

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The gaseous carbonaceous material may be any carbonaceous compound or substance which may be gasified and which is capable of forming carbon nanotubes when subjected to pyrolysis. Examples of such compounds are alkanes, alkenes, alkynes and aromatic hydrocarbons, for example, methane, ethylene, benzene or acetylene. Preferably the carbonaceous material is acetylene.

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Pyrolysis is performed in the presence of a catalyst. The catalyst may be any compound, element or substance suitable for catalysing the conversion of a carbonaceous material to

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carbon nanotubes under the pyrolysis conditions. Preferably the catalyst comprises a transition metal including Ni, Fe, Co, Al, Mn, Pd, Cr or alloys thereof in any suitable oxidation state. Most preferably, the catalyst comprises Ni. For example, the catalyst may be prepared from polyvinylalcohol/ $\text{Ni}(\text{NO}_3)_2 \cdot 6\text{H}_2\text{O}$ (PVA Ni.) Preferably, the surface of
5 the substrate is coated with a substance from which the catalyst is prepared. For example, a spin-coated PVA Ni layer, subjected to oxidation at 500°C for 30 minutes and reduction at 600°C for 30 minutes provided a catalyst coating showing strong adhesion onto a glass substrate, even when subjected to compressed air. Reduction of the coating in the reactor is readily performed by supplying a mixture of H_2/Ar to provide the catalyst-coated
10 substrate. The substrate may then be maintained in an inert atmosphere, for example, nitrogen or argon, to prevent the catalyst being oxidised.

The pyrolysis conditions employed will depend on the nature of the gaseous carbonaceous material, the catalyst used, and the length and density of the carbon nanotubes required. It
15 is possible to vary the pyrolysis conditions, such as temperature, time, catalyst, pressure or flow rate through the reactor to obtain carbon nanotubes having different characteristics.

Pyrolysis may be performed at temperatures above 300°C . Preferably in the process of the invention temperatures in the heating zone are between 400°C and 800°C . The selection
20 of catalyst affects the temperature at which carbon nanotubes may be formed. The carbon formed during pyrolysis is then selectively deposited on the hot surface of the substrate in the heating zone, forming carbon nanotubes. Temperatures below 400°C are demonstrated to be suitable for the nanotube growth with the ratio of carbon nanotubes to carbon nanofibre, their morphology and alignment depending on the conditions used.
25 Surprisingly, it was found using the process of the present invention that well-aligned carbon nanotubes were easily formed well below the softening point of normal glass plates (*ca.* 640°C). Within the temperature ranging from 400°C and 800°C , the nanotube deposition is completed within a couple of seconds to 20 minutes. The formation of carbon nanotubes is a typical transient reaction and its deposition rate can be controlled by
30 adjusting the pressure of the carbonaceous gas. With a low feed of carbonaceous gas, the carbon nanotube growth is from edge to centre of the substrate. In contrast, a homogenous

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coating is seen over the substrate surface within a couple of seconds at a high gas feed rate. Therefore, the deposition reaction can be region-specifically controlled by controlling the feed of the carbonaceous gas.

- 5 The carbon nanotubes produced by the process may be aligned or non-aligned. Aligned or non-aligned carbon nanotubes may be selected for by varying temperatures, type of catalyst used and the density of the catalyst coating on the substrate. For example, a low density of catalyst coating will favour non-aligned carbon nanotube growth whereas a high density of catalyst coating will favour aligned carbon nanotube growth.

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The length of aligned carbon nanotubes may be varied over a certain range (from a sub-micrometre to several tens of micrometres) in a controllable fashion by changing the experimental conditions such as the pyrolysis time and gas flow rate. The size and shape of the aligned carbon nanotube film is, in principle, limited only by the size and shape of the substrate.

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According to one embodiment of the invention, the reaction chamber may have a pre-heating zone, where the substrate may be pre-heated to a predetermined temperature before entering the localised heating zone where carbon nanotube deposition occurs. The reaction chamber may also have a cooling zone where the substrate is cooled after carbon nanotube deposition is complete. In one embodiment of the invention, the reactor includes a means of moving a substrate from the pre-heating zone to the localised heating zone and from the localised heating zone to the cooling zone. For example, the substrate may sit on a transporting belt.

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The reaction chamber of the present invention may be adapted to have multiple localised heating zones by having multiple heating elements located within the chamber. Carbon nanotubes may then be deposited on multiple substrates simultaneously or in separate areas of a larger substrate. In a reactor containing multiple localised heating zones, a gas inlet may be attached to a gas distributor which allows the gaseous carbonaceous material to be supplied to each of the multiple localised heating zones simultaneously. Alternatively, the

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gaseous carbonaceous materials may be supplied through an inlet at one end of the chamber and allowed to flow across each substrate and through each localised heating zone sequentially.

- 5 The reactor of the present invention may be adapted to allow continuous carbon nanotube deposition on multiple substrates by having a means of moving a substrate from a pre-heating zone to the localised heating zone before carbon nanotube deposition and a means of moving a substrate to a cooling zone after carbon nanotube deposition is complete. Continuous carbon nanotube production may be achieved by pre-heating a substrate or
- 10 substrates in the pre-heating zone, moving the substrate or substrates to the localised heating zone, synthesising carbon nanotubes on the substrate or substrates, moving the substrate or substrates into the cooling zone, removing the cooled substrate or substrates having a carbon nanotube film deposited on it/them from the reactor, and supplying new
- 15 substrate or substrates to the pre-heating zone. Generally only one step, e.g. pre-heating, carbon deposition or cooling, is performed at a certain part of the reactor at any one time. The process may be continuously repeated. The reduction of a substance from which the catalyst is prepared may also be performed in the reactor by supplying a mixture of Ar/H₂ to the chamber to provide the catalyst coated substrate.
- 20 In the continuous process, the substrate may be located in a first chamber and then transported to a second chamber having a pre-heating zone and/or localised heating zone and/or cooling zone. Advantageously, the reduction of a substance from which the catalyst is prepared may be performed in the first chamber to provide a catalyst coated substrate. The substrate may then be transported to the pre-heating zone or localised heating zone in
- 25 the second chamber when the atmosphere in the first chamber has been flushed with inert gas. In a preferred embodiment, the two chambers are connected and the catalyst coated substrate may pass from the first chamber to the pre-heating zone or localised heating zone in the second chamber through a connecting door.
- 30 In a reactor adapted for use in the continuous process of the invention, the localised heating zone and the cooling zone may be in the same chamber. Alternatively, the cooling

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zone may be in a separate chamber from the localised heating zone. If the localised heating zone and cooling zone are in separate chambers, the two chambers may be connected such that the substrate in the localised heating zone may be transferred to the cooling zone via a connecting door after carbon deposition has occurred.

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According to the present invention it is possible to prepare multilayer carbon nanotube materials by synthesising a first layer of carbon nanotubes on a substrate under a first set of pyrolysis conditions, and then synthesising a second layer of carbon nanotubes on the nanotube coated substrate under a second set of pyrolysis conditions.

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This process may be repeated until the desired number of carbon nanotube layers are present. Each layer of carbon nanotubes may be deposited using the same or different pyrolysis conditions. After preparation of the multilayer structure the carbon nanotube film may be removed from the substrate using appropriate conditions.

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It is also possible according to the present invention to prepare hetero-structured multilayer carbon nanotube films by interposing layers of carbon nanotubes between layers of pyrolysis resistant materials, the carbon nanotubes being generated in accordance with the process of the present invention.

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The term "hetero-structured" as used herein refers to a multilayer structure which includes one or more carbon nanotube layers together with layers of other materials

The pyrolysis resistant material may be a metal, preferably Au, Pt, Ni, Cu, a semiconductor, TiO₂, MgO, Al₂O₃, ZnO, SnO₂, Ga₂O₃, In₂O₃, CdO or a polymer or any other pyrolysis resistant material that is capable of supporting the nanotube growth.

The pyrolysis resistant material may be applied to the carbon nanotube coated substrate by any suitable means. Preferably metals are applied by sputter-coating, polymers are applied by spin-casting and semiconductors by sputter-coating or physical deposition.

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As is evident from the above description, the process and apparatus of the invention allow the preparation of a large variety of carbon nanotube films and structures. It is also possible to provide patterned layers using appropriate masking and etching techniques. At lower temperatures it may be possible to use the reactor of the invention to prepare carbon
5 nanofibres or mixtures of carbon nanotubes and nanofibres.

The materials produced by the present invention may be used in the construction of devices for practical applications in many fields including electron emitters, field-emission transistors, electrodes for photovoltaic cells and light emitting diodes, optoelectronic
10 elements, bismuth actuators, chemical and biological sensors, gas and energy storages, molecular filtration membranes and energy-absorbing materials.

The invention can be more fully understood from the following detailed description of Figure 1 and the examples. It should be understood that the examples and Figures
15 described are only for illustration purposes, which does not intend to constitute a limitation on the invention.

Brief Description of the Drawings

20 Figure 1AA is a diagrammatic side-view representation of a pyrolysis flow reactor of the invention having a gas inlet at one end of the chamber, and a gas outlet at the other.

Figure 1AB is a diagrammatic side view representation of a pyrolysis flow reactor of the invention having a gas inlet positioned above the localised heating zone and gas outlets
25 positioned on the respective ends of the reaction chamber.

Figure 1AC is a diagrammatic end view representation of a pyrolysis reactor of the invention having a gas inlet positioned above the localised heating zone and a gas outlet positioned below.
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Figure 1BA is a diagrammatic side view representation of a pyrolysis reactor of the

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invention having multiple localised heating zones.

Figure 1BB is a diagrammatic top view representation of a heating element supporting multiple substrates.

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Figure 1CA is a diagrammatic side view representation of a pyrolysis reactor of the invention having a pre-heating zone, a localised heating zone and a cooling zone and wherein inlet for supplying the gaseous carbonaceous material is positioned above the localised heating zone.

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Figure 1CB is a diagrammatic top plan view of a heating element having a pre-heating zone, a localised heating zone and a cooling zone.

Figure 1D is a diagrammatic side plan view of a reactor of the invention in which a substrate may be moved from the pre-heating zone to the localised heating zone and finally to the cooling zone before removal from the reactor.

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Figure 1EA is a series of diagrammatic side plan views of a heating element designs suitable for use in the reactor of the invention.

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Figure 1EB is a diagrammatic top plan view of a heating element suitable for use in the reactor of the invention.

Figure 1AA is a diagrammatic representation of a pyrolysis reactor of the invention in which a substrate (1) has been positioned on the heating element (2). The heating element (2) and substrate (1) are located within a reaction chamber (7) defined by chamber walls (3). A sheet of conducting material (4) is positioned between the substrate (1) and the heating element (2) in order to provide homogenous heating of the substrate (1). A gas inlet (5) is positioned at one end of the reaction chamber and a gas outlet (6) is positioned at the opposing end of the reaction chamber such that any gas introduced into the reaction chamber (7) flows from one end of the chamber to the other, flowing over the substrate (1).

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Figure 1AB is a diagrammatic representation of a pyrolysis reactor of the invention similar to that shown in Figure 1AA but having a gas inlet (5) positioned above the heating element (2) upon which the substrate (1) is supported. The gas is supplied directly into the
5 localised heating zone (8). Gas outlets (6) are located at each end of the chamber.

Figure 1AC is a diagrammatic end view representation of a pyrolysis reactor of the invention similar to that shown in Figures 1AA or 1AB but having a gas outlet (6) located below the localised heating zone (8).

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In operation, the reactor shown in Figures 1AA to 1AC is flushed with an inert gas such as N₂ or Ar, or is evacuated by means of a vacuum pump connected to the gas outlet (6). A mixture of H₂ and Ar is introduced into the reaction chamber (7) to ensure the catalyst is in a reduced state. The heating element (2) is heated to the required temperature which heats
15 the substrate (1) to the required temperature. When the catalyst has been reduced, a flow of Ar gas is maintained to ensure an inert atmosphere in the chamber. A gaseous carbonaceous material, such as ethylene, is introduced into the reaction chamber (7) through the gas inlet (5). Pyrolysis of the carbonaceous material occurs in the localised heating zone (8) and carbon is deposited on the hot surface of the substrate (1) to produce a
20 carbon nanotube layer. The substrate (1) is then allowed to cool.

Figure 1BA is a diagrammatic side view representation of a pyrolysis reactor of the invention in which multiple substrates (1) are positioned on a heating element (2). The heating element (2) and the substrates are located within a reaction chamber (7) defined by
25 chamber walls (3). The gas inlet (5) is connected to a gas distributor (9) that allows the carbon-containing material to be simultaneously introduced into multiple localised heating zones (8). The gas outlet (6) is positioned at one end of chamber (7) and a connection to a vacuum pump (10) is positioned at the opposing end of chamber (7).

30 Figure 1BB is a diagrammatic top plan view representation of the heating element (2) of the reactor shown in Figure 1BA. Multiple substrates (1) are positioned on the heating

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element (2).

In operation, the reactor shown in Figure 1BA is evacuated by a vacuum pump connected to outlet (10) and a mixture of H₂ and Ar is introduced into the reaction chamber (7). The heating element (2) is heated to the required temperature which heats the multiple substrates (1) to the required temperature. When the catalyst has been reduced, a positive pressure of inert atmosphere (Ar) is maintained in the reaction chamber (7). A gaseous carbonaceous material is introduced into multiple localised heating zones (8) from the gas inlet (5) by means of a gas distributor (9). Pyrolysis of the carbonaceous material occurs in the multiple localised heating zones (8) and carbon is deposited on the hot surfaces of the multiple substrates (1) to produce a carbon nanotube layer on each of the multiple substrates (1). The substrates are then allowed to cool. Multiple substrates are simultaneously coated with a layer of carbon nanotube in this reactor.

Figure 1CA is a diagrammatic side view representation of a pyrolysis reactor of the invention in which a substrate (1) is positioned on the heating element (2). The heating element (2) and the substrate (1) is located within the reaction chamber (7) defined by chamber walls (3). The heating element (2) is divided into zones having different temperatures, a pre-heating zone (11), a localised heating zone (8) and a cooling zone (12). The gas inlet (5) is positioned above the localised heating zone (8) and distributes the gaseous carbonaceous material evenly over the substrate (1) when it is in the localised heating zone (8). The gas outlet (6) is positioned at one end of the chamber (7) and a connection to a vacuum pump (10) is positioned at the opposing end of the chamber (7).

Figure 1CB is a diagrammatic top plan view representation of a heating element (2) of the reactor shown in Figure 1CA. A substrate (1) is initially positioned in the pre-heating zone (11), then the substrate is moved to the localised heating zone and finally the substrate is moved to the cooling zone (12).

In operation, a substrate (1) is pre-heated to a predetermined temperature in the pre-heating zone (11). The substrate (1) is then moved to the localised heating zone (8) and is heated

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to pyrolysis temperature. A gaseous carbonaceous material is introduced into the localised heating zone (8) through gas inlet. The gaseous carbonaceous material is pyrolysed and carbon nanotubes are deposited on the surface of the substrate (1). The substrate (1), upon which carbon nanotubes have been deposited, is moved to the cooling zone (12) and
5 allowed to cool before being removed from the reactor. The reactor may also include a mechanism for transferring the cooled substrate (1) having a film of carbon nanotubes out of the reactor and a mechanism for positioning a new substrate in the pre-heating zone. The reactor may also include a mechanism which allows transfer of a substrate from one zone to the next within the chamber. (7) The process may be performed in a continuous
10 manner, for example, the substrate may sit on a transporting belt. Such a device is also illustrated in Figure 1D.

Figures 1EA to 1ED show a series of heating element (2)/conducting material (4)/substrate (1) configurations which may be useful in the reactor of the invention. In Figure 1EA, the
15 sheet of conducting material (4) and the substrate (1) are embedded in the heating element (2). In Figure 1EB, the sheet of conducting material (4) is embedded in the heating element (2) and a substrate (1) which is smaller than the heating element (2) is placed in the heating element (2) on the conducting metal sheet (4). In Figure 1EC, the surface of the heating element (2) is flat and a substrate (1) having a smaller surface than the heating
20 element (2) is placed on a sheet of conducting material (4) also having a smaller surface than the heating element (2). The sheet of conducting material (4) and the substrate (1) protrude above the surface of the heating element (2). In Figure 1ED, the heating element (2) is coated with a conducting material (4) across its entire surface and a substrate (1) is positioned on top of the conducting material coating. The configuration in Figure 1ED is
25 the most suitable for use in a reactor where continuous carbon nanotube deposition occurs, for example, the reactor shown in Figure 1C or Figure 1D.

Figure 1EE is a diagrammatic top plan view of a preferred heating element (2) having a substrate (1) positioned on it. The heating element (2) has a sheet of conducting material
30 (4) placed on it to provide homogenous temperature distribution and a resistant wire (13) is supported within the heating element to heat the heating element.

Examples

Example 1: Preparation of non-aligned carbon nanotubes

5 A glass substrate was spin-coated with a PVA Ni layer (100mM $\text{Ni}(\text{NO}_3)_2 \cdot 6\text{H}_2\text{O}$ and 3 wt% PVA) to provide a catalyst for deposition of carbon nanotubes. The coated substrate was oxidised at 500°C for 30 minutes and reduced at 600°C for 30 minutes. After such treatment, the catalyst coating was strongly adhered to the substrate. The catalyst-coated substrate was placed in the reactor as shown in Figure 1A and the atmosphere in the
10 reaction chamber was replaced with H_2/Ar to ensure the catalyst was in a reduced state. The substrate was heated at 650°C on the heating element and acetylene/Ar gas (V:V = 1:3), introduced at a total flow rate of 60 ml/min for a time of 3 minutes, was pyrolysed resulting in the deposition of non-aligned carbon nanotubes on the catalyst-coated substrate.

15

Example 2: Preparation of aligned carbon nanotubes

Aligned carbon nanotubes were prepared by the same method as applied in Example 1, but at a lower temperature (440°C) was used. The resulting carbon nanotubes align almost normal to the substrates surface and are densely packed with a fairly uniform tubular
20 length of *ca.* 1 μm .

THE CLAIMS DEFINING THE INVENTION ARE AS FOLLOWS:

1. 1. A process for preparing carbon nanotubes comprising:
locating a substrate capable of supporting carbon nanotube growth in a localised
5 heating zone within a reaction chamber, said localised heating zone being provided
by a heating element located within said reaction chamber,
passing a gaseous carbonaceous material into said reaction chamber such that the
gaseous material passes over and contacts said substrate in the localised heating
zone, whereby said gaseous material undergoes pyrolysis under the influence of
10 said heat to form carbon nanotubes on said substrate.
2. A process according to claim 1 wherein the localised heating zone has a
temperature greater than 300°C.
3. A process according to claim 2 wherein the localised heating zone has a
temperature between 400°C and 800°C.
- 15 4. A process according to claim 1 wherein the substrate is quartz glass, mesoporous
silica, nanoporous alumina, a ceramic plate, glass, graphite or mica.
5. A process according to claim 4 wherein the substrate is glass.
6. A process according to claim 1 wherein the gaseous carbonaceous material is
selected from an alkane, alkene, alkyne or aromatic hydrocarbon.
- 20 7. A process according to claim 6 wherein the gaseous carbonaceous material is
selected from methane, ethylene, benzene or acetylene.
8. A process according to claim 7 wherein the gaseous carbonaceous material is
acetylene.
9. A process according to claim 1 wherein pyrolysis of the carbonaceous material
25 occurs in the presence of a catalyst.
10. A process according to claim 9 wherein the catalyst is coated on the substrate.
11. A process according to claim 9 or 10 wherein the catalyst comprises a transition
metal selected from Ni, Fe, Co, Al, Mn, Pd, Cr or alloys thereof.
12. A process according to claim 11 wherein the catalyst comprises Ni.
- 30 13. A process according to claim 1 wherein the pyrolysis conditions are controlled to
provide aligned carbon nanotubes.

- 18 -

14. A process according to claim 1 wherein the pyrolysis conditions are controlled to provide non-aligned carbon nanotubes.
15. A process according to claim 1 wherein the pyrolysis conditions are controlled to provide homogeneous carbon nanotube growth on the substrate.
- 5 16. A process according to claim 1 wherein the pyrolysis conditions are controlled to provide patterned carbon nanotube growth on the substrate.
17. A process for preparing multilayer carbon nanotube materials comprising:
 - (a) synthesising a first layer of carbon nanotubes on a substrate under a first set of pyrolysis conditions to provide a nanotube coated substrate;
 - 10 (b) synthesising a second layer of carbon nanotubes on the nanotube coated substrate under a second set of pyrolysis conditions, wherein at least one of steps (a) and (b) is performed using a process of claim 1.
18. A process according to claim 17 wherein step (b) is repeated at least once.
19. A process according to claim 17 wherein the pyrolysis conditions of step (a) are the same as the pyrolysis conditions of step (b).
- 15 20. A process according to claim 17 wherein the pyrolysis conditions of step (a) are different from the pyrolysis conditions of step (b).
21. A process for the preparation of a hetero-structured multilayer carbon nanotube film comprising:
 - 20 (a) synthesising a first layer of carbon nanotubes on a substrate under a first set of pyrolysis conditions to provide a nanotube coated substrate;
 - (b) coating a layer of pyrolysis resistant material onto the nanotube coated substrate to provide a hetero-structured multilayer substrate;
 - (c) synthesising a second layer of carbon nanotubes on the hetero-structured multilayer substrate under a second set of pyrolysis conditions,
 - 25 wherein at least one of steps (a) and (c) is performed using a process of claim 1.
22. A process according to claim 21 wherein steps (b) and (c) are repeated at least once.
23. A process according to claim 21 wherein the pyrolysis conditions of step (a) are the same as the pyrolysis conditions of step (c).
- 30

- 19 -

24. A process according to claim 21 wherein the pyrolysis conditions of step (a) are different from the pyrolysis conditions of step (c).
25. A process according to claim 21 wherein the pyrolysis resistant material is a metal, a semiconductor or a polymer.
- 5 26. A process according to claim 25 wherein the pyrolysis resistant material is a metal.
27. A reactor for preparing carbon nanotubes comprising:
a reaction chamber,
at least one support means located within said reaction chamber capable of supporting a substrate, said substrate being capable of supporting carbon nanotube growth,
10 at least one heating element located within said reaction chamber capable of providing localised heating to said substrate within said reaction chamber,
means for passing a gaseous carbonaceous material into said reaction chamber such that it passes over and contacts said substrate.
- 15 28. A reactor according to claim 27 wherein the reaction chamber is formed from metal, glass, plastic or a combination thereof.
29. A reactor according to claim 28 wherein the reaction chamber is formed from glass or comprises at least one glass panel.
30. A reactor according to claim 27 wherein the heating element comprises resistant wires, an induction field, microwave radiation or infrared radiation.
20 31. A reactor according to claim 27 wherein the heating element is located within the substrate support.
32. A reactor according to claim 31 wherein the heating element and substrate support comprise a ceramic plate into which resistant wires have been inserted.
- 25 33. A reactor according to claim 27 wherein the means for passing a gaseous carbonaceous material into the reaction chamber is at least one gas conduit.
34. A reactor according to claim 33 wherein the at least one gas conduit is located above the substrate.
35. A reactor according to claim 33 wherein the at least one gas conduit is located to
30 allow the gaseous carbonaceous material to flow across the surface of the substrate.
36. A reactor according to claim 27 comprising multiple support means.

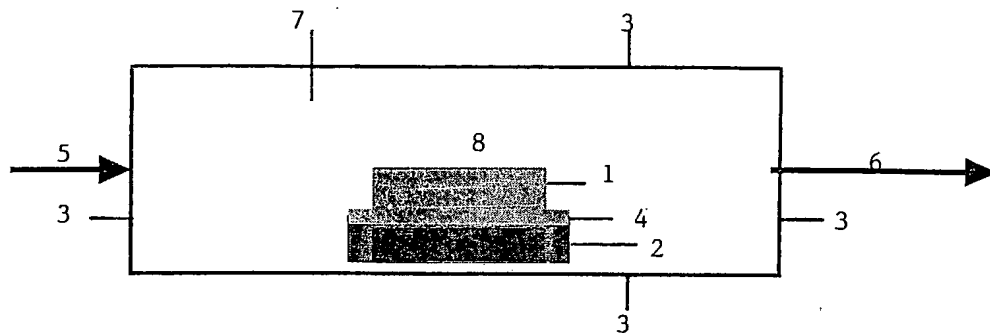
- 20 -

37. A reactor according to claim 27 comprising multiple heating elements.
38. A reactor according to claim 27 further comprising a pre-heating zone.
39. A reactor according to claim 38 wherein the pre-heating zone is located in a separate chamber from the reaction chamber.
- 5 40. A reactor according to claim 27 further comprising a cooling zone.
41. A reactor according to claim 40 wherein the cooling zone is located in a separate chamber from the reaction chamber.
42. A reactor according to claim 27 further comprising a means of transferring a substrate from a pre-heating zone to a support means and/or from the support means to a cooling zone.
- 10
43. Carbon nanotubes prepared by the process of any one of claims 1 to 16.
44. Multilayer carbon nanotubes prepared by the process of any one of claims 17 to 20.
45. Hetero-structured multilayer carbon nanotube films prepared by the process of any one of claims 21 to 26.

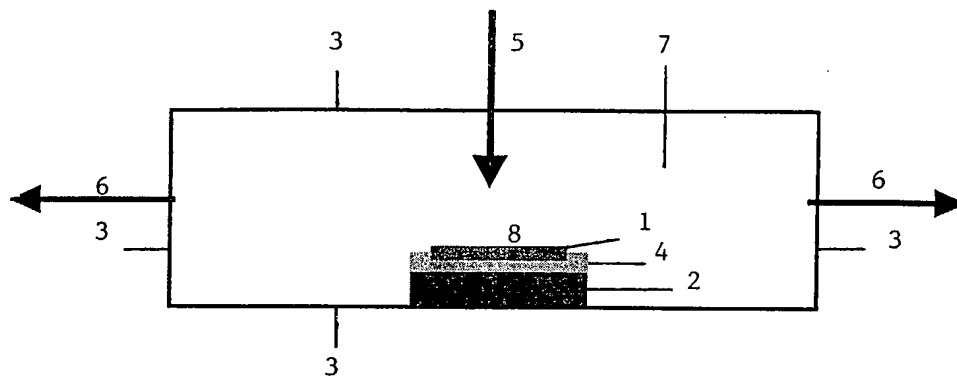
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Figure 1A

A,



B,



C,

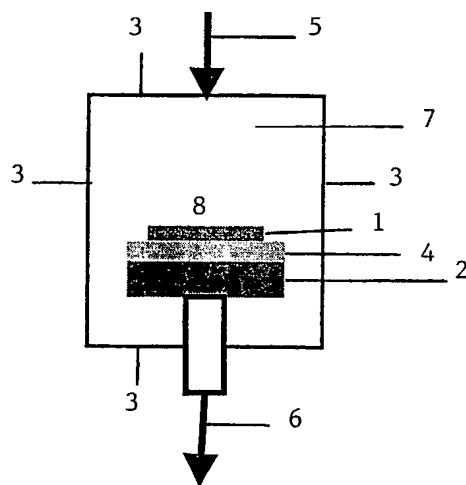
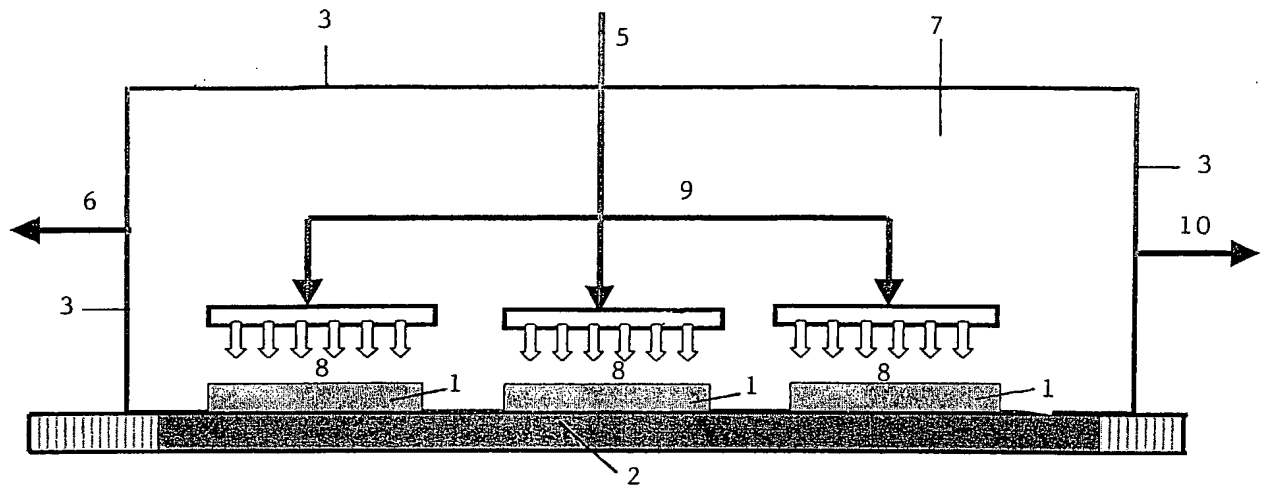


Figure 1B.

A



B

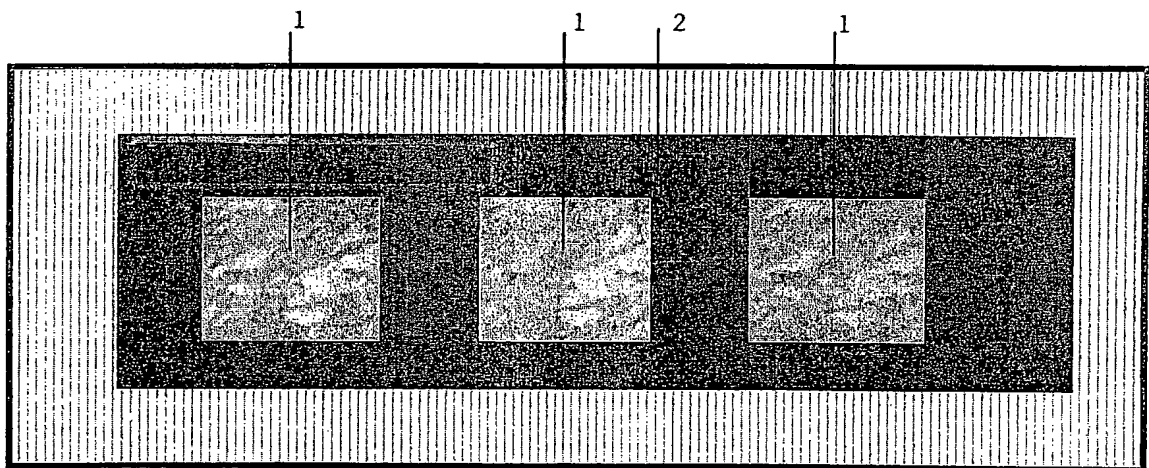
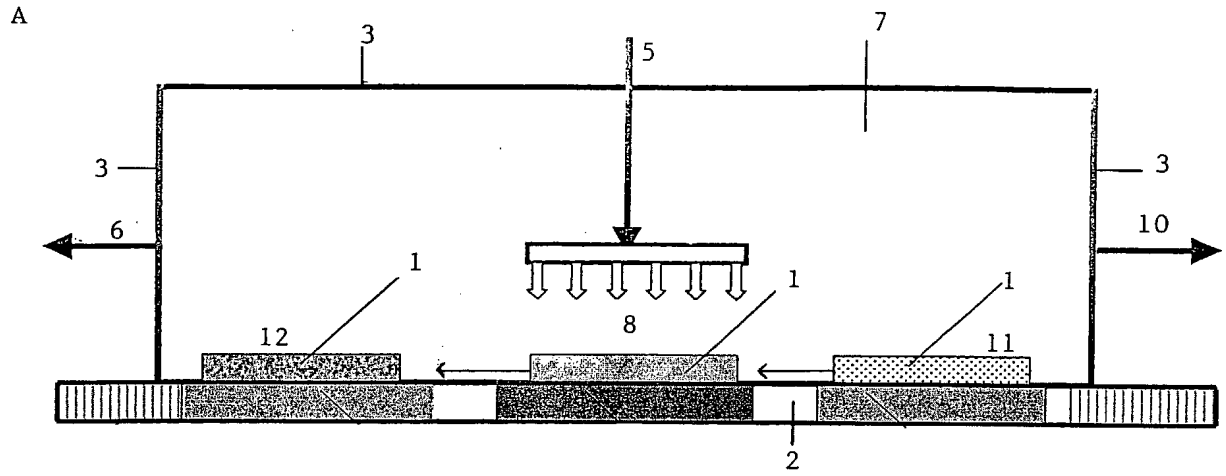


Figure 1C



B

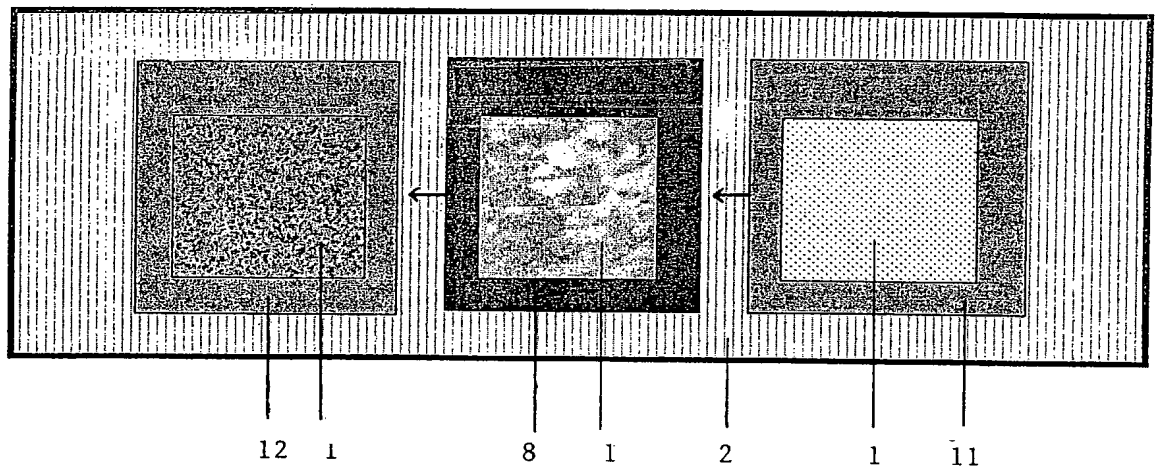


Figure 1 D

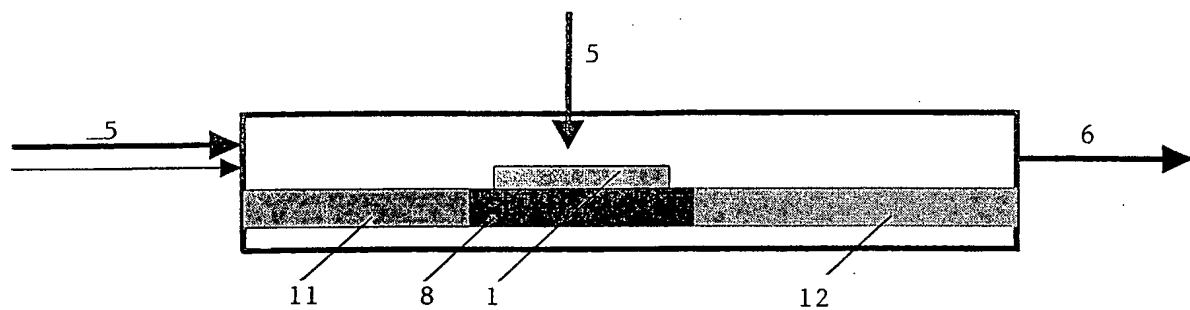
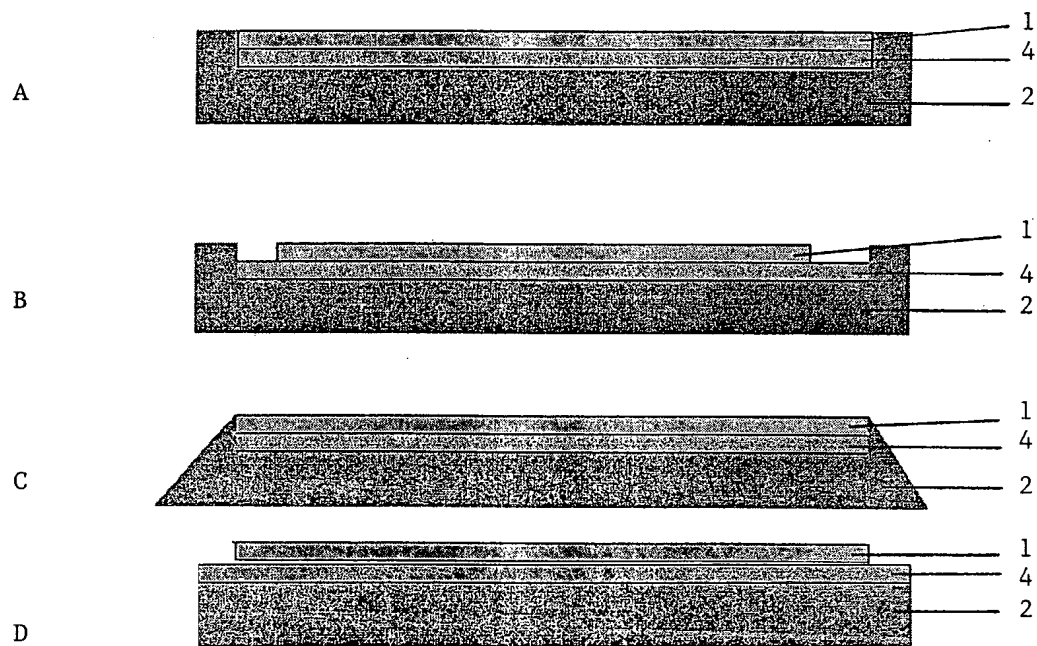
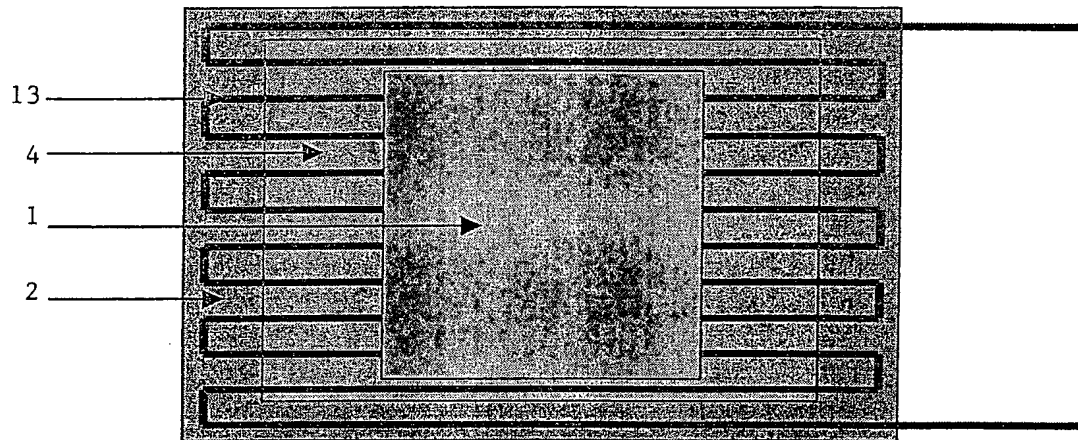


Figure 1E



E



INTERNATIONAL SEARCH REPORT

International application No.

PCT/AU02/00437

A. CLASSIFICATION OF SUBJECT MATTER												
<i>Int.Cl.</i> ⁷ : B82B 3/00, 1/00, C01B 31/02, C23C 16/46, D01F 9/12, 9/127, 9/133												
According to International Patent Classification (IPC) or to both national classification and IPC												
B. FIELDS SEARCHED												
Minimum documentation searched (classification system followed by classification symbols)												
<i>Int.Cl.</i> ⁷ : B82B 3/00, 1/00, C01B 31/02, D01F 9/12, 9/127, 9/133, C23C 16/46, 16/26												
Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched												
Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)												
DWPI: <i>Int.Cl.</i> ⁷ as above, nanotube+, nano+ and tube?, carbon, fuller+, buckmin+, c60, c70, heat+, temperature?, substrate?, holder, local+, reactor, reaction chamber, furnace, pyrolysis												
C. DOCUMENTS CONSIDERED TO BE RELEVANT												
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.										
A	WO 00/73204 A1 (COMMONWEALTH SCIENTIFIC AND INDUSTRIAL RESEARCH ORGANISATION) 7 December 2000 Whole document											
A	WO 00/73203 A1 (COMMONWEALTH SCIENTIFIC AND INDUSTRIAL RESEARCH ORGANISATION) 7 December 2000 Whole document											
A	WO 00/63115 A1 (COMMONWEALTH SCIENTIFIC AND INDUSTRIAL RESEARCH ORGANISATION) 26 October 2000 Whole document											
<input checked="" type="checkbox"/> Further documents are listed in the continuation of Box C <input checked="" type="checkbox"/> See patent family annex												
<p>* Special categories of cited documents:</p> <table border="0"> <tr> <td>"A" document defining the general state of the art which is not considered to be of particular relevance</td> <td>"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention</td> </tr> <tr> <td>"E" earlier application or patent but published on or after the international filing date</td> <td>"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone</td> </tr> <tr> <td>"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)</td> <td>"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art</td> </tr> <tr> <td>"O" document referring to an oral disclosure, use, exhibition or other means</td> <td>"&" document member of the same patent family</td> </tr> <tr> <td>"P" document published prior to the international filing date but later than the priority date claimed</td> <td></td> </tr> </table>			"A" document defining the general state of the art which is not considered to be of particular relevance	"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention	"E" earlier application or patent but published on or after the international filing date	"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone	"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art	"O" document referring to an oral disclosure, use, exhibition or other means	"&" document member of the same patent family	"P" document published prior to the international filing date but later than the priority date claimed	
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"E" earlier application or patent but published on or after the international filing date	"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone											
"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art											
"O" document referring to an oral disclosure, use, exhibition or other means	"&" document member of the same patent family											
"P" document published prior to the international filing date but later than the priority date claimed												
Date of the actual completion of the international search 3 May 2002		Date of mailing of the international search report 17 MAY 2002										
Name and mailing address of the ISA/AU -AUSTRALIAN PATENT OFFICE PO BOX 200, WODEN ACT 2606, AUSTRALIA E-mail address: pct@ipaustalia.gov.au Facsimile No. (02) 6285 3929		Authorized officer RAJEEV DESHMUKH Telephone No : (02) 6283 2145										

INTERNATIONAL SEARCH REPORT

International application No.

PCT/AU02/00437

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	EP 1061041 A1 (ILJIN NANOTECH CO., LTD.) 20 December 2000 Column 4, line 53 - column 5, line 10	
	WO 00/76912 A2 (BATTELLE MEMORIAL INSTITUTE) 21 December 2000	
A	Page 5, lines 16-28	
A	WO 01/21863 A1 (COMMONWEALTH SCIENTIFIC AND INDUSTRIAL RESEARCH ORGANISATION) 20 March 2001 Whole document	
	US 2001/0009693 A1 (LEE et al.) 26 July 2001	
P, X	Page 2, left column, line 49 - right column, line 3	1,27
P, X	WO 01/23303 A1 (ELECTROVAC, FABRIKATION ELEKTROTECHNISCHER SPEZIALARTIKEL GESELLSCHAFT M.B.H. et al.) 5 April 2001 Abstract; page 5, line 32 - page 6, line 5; figure 1	1,27

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No.

PCT/AU02/00437

This Annex lists the known "A" publication level patent family members relating to the patent documents cited in the above-mentioned international search report. The Australian Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

Patent Document Cited in Search Report			Patent Family Member		
WO	200073204	AU	200045284	EP	1198414
WO	200073203	AU	200045283	EP	1200341
WO	200063115	AU	200036496	EP	1183210
EP	1061041	JP	2001032071	CN	1278021
WO	200076912	AU	200078244	US	2002004136
WO	200121863	AU	200076340	US	6361861
US	2001009693	CN	1315588	EP	1149932
WO	200123303	AT	1667/99	JP	2001234341
AU				AU	200065483
END OF ANNEX					